AMENDMENT TO THE CLAIMS:

This listing of claims will replace all prior versions, and listings of claims in the application:

LISTING OF CLAIMS:

The list of currently pending claims is presented below.

WHAT IS CLAIMED IS:

1. (Original) A xanthene dye having the formula:

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in which

R¹, R², R⁴, R⁵, R⁶, R⁷, R⁸, R⁹, R¹⁰ and R¹¹ are independently selected from substituted or unsubstituted alkyl, substituted or unsubstituted heteroalkyl, substituted or unsubstituted aryl, substituted or unsubstituted heteroaryl, substituted or unsubstituted heterocycloalkyl, halogen, H, NO₂, CN and C(Z¹)R¹⁴, NR¹⁵R¹⁶ and Z²R¹⁶;

9 R^3 is selected from Z^2R^{16} and $NR^{15}R^{16}$

10	wherein
11	Z ¹ is a member selected from O, S and NH;
12	Z^2 is a member selected from O and S;
13	R ¹⁵ is a member selected from H, substituted or unsubstituted alkyl, and
14	substituted or unsubstituted heteroalkyl;
15	R ¹⁶ is selected from H, substituted or unsubstituted alkyl, substituted or
16	unsubstituted heteroalkyl, C(Z ³)R ¹⁷ , and a nitrogen-containing reactive
17	group comprising R ¹⁵ and R ¹⁶ , together with the nitrogen to which they are
18	attached, wherein said reactive group is a member selected from -NHNH ₂ ,
19	-N=C=S and $-N=C=O$
20	wherein
21	Z ³ is a member selected from O, S and NH;
22	R ¹⁷ is a member selected from substituted or unsubstituted alkyl,
23	substituted or unsubstituted heteroalkyl, OR ¹⁸ , and NR ¹⁹ R ²⁰
24	wherein
25	R ¹⁸ is a member selected from H, substituted or unsubstituted
26	alkyl, substituted or unsubstituted heteroalkyl, substituted
27	or unsubstituted aryl, substituted or unsubstituted
28	heteroaryl and C(O)R ²¹
29	wherein
30	R ²¹ is substituted or unsubstituted alkyl or substituted or
31	unsubstituted heteroalkyl;
32	R ¹⁹ and R ²⁰ are members independently selected from H,
33	substituted or unsubstituted alkyl and substituted or
34	unsubstituted heteroalkyl
35	Y is a member selected from $C(O)$ and $S(O)_2$;

36	X is a member selected from (NR ²² R ²³) and (O)	
37	wherein	
38 39	R ²² and R ²³ are members independently selected from H, substituted or unsubstituted alkyl and substituted or unsubstituted heteroalkyl; and	
10	R ¹² and R ¹³ are members independently selected from substituted or unsubstituted alkyl,	
11	substituted or unsubstituted heteroalkyl, substituted or unsubstituted	
12	heterocycloalkyl, substituted or unsubstituted aryl and substituted or unsubstitute	d
13	heteroaryl, with the proviso that at least one of R ¹² or R ¹³ comprises a member	
14	selected from a bond to a carrier molecule, a bond to a linker bound to a carrier	
15	molecule, a bond to a solid support, a bond to a linker attached to a solid support,	,
16	a bond to a fluorescence quencher, a bond to a linker to a fluorescence quencher	
17	and an oxygen-containing reactive group, and further with the proviso that when	
18	R ¹² and R ¹³ , together with the nitrogen to which they are attached form a	
19	piperazine ring said oxygen-containing reactive group is a phosphoramidite and	
50	said bond to a carrier molecule is other than a bond to a peptide.	
1	2. (Original) The xanthene dye according to claim 1, wherein R ³ is R ¹⁵ R ¹⁶ N; and X	r L
2	is NR ²³ R ²⁴ , wherein R ¹⁵ , R ¹⁶ , R ²³ and R ²⁴ are members independently selected from H,	
3	substituted or unsubstituted alkyl and substituted or unsubstituted heteroalkyl.	
1	3. (Original) The xanthene dye according to claim 1, wherein at least one of R ⁸ , R ⁹	,
2	R ¹⁰ and R ¹¹ is a halogen.	
1	4. (Original) The xanthene dye according to claim 1, wherein R ⁹ and R ¹⁰ are	
2	halogen.	
1	5. (Original) The xanthene dye according to claim 3, wherein R ³ is OR ¹⁶ ; and X is	
2	O.	
1	6. (Original) The xanthene dye according to claim 5, wherein R ² and R ⁶ are	
2	halogen.	

7. (Original) The xanthene dye according to claim 5, wherein R² and R⁶ are independently selected from substituted or unsubstituted alkyl and substituted or unsubstituted heteroalkyl.

8. (Original) The xanthene dye according to claim **1**, wherein R³ is NR¹⁵R¹⁶ and R², R⁴ and R¹⁵ and R¹⁶, together with the nitrogen atom to which they are bound, are fused with the phenyl moiety to which NR¹⁵R¹⁶, R² and R⁴ are bound, forming a substituted or unsubstituted ring system having formula:

9. (Original) The xanthene dye according to claim 1, wherein X is NR²²R²³ and R⁵, R⁶ and R²² and R²³, together with the nitrogen atom to which they are bound, are fused with the unsaturated 6-member ring to which NR²²R²³, R⁵ and R⁶ are bound, forming a substituted or unsubstituted ring system having the formula:

10. (Original) The xanthene dye according to claim 1, wherein said oxygen-containing reactive functional group is a member selected from hydroxyl and activated derivatives thereof, phosphoramidite, and carboxylic acid and activated derivatives thereof.

11. (Original) The xanthene dye according to claim 1, wherein R¹² and R¹³, together with the nitrogen to which they are bound are joined to form a ring system.

1 12. (Original) The xanthene dye according to claim 11, wherein NR¹²R¹³ has the 2 formula:

$$\xi$$
 R^{25}

34 wherein

h and i are members independently selected from integers such that the sum (h + i) is

6 from 4-8; and

 R^{25} is a reactive functional group.

1 13. (Original) The xanthene dye according to claim 1, wherein R¹² comprises a 2 moiety having the formula:

$$\xi - O - L^1 - L^2 - L^3 - O + \xi$$

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5 L¹, L² and L³ are members independently selected from substituted or unsubstituted alkyl and substituted or unsubstituted heteroalkyl; and

7 t is 0 or 1.

1 14. (Original) The xanthene dye according to claim 13, said moiety having the 2 formula:

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R^x and R^y are members independently selected from H, substituted or unsubstituted alkyl, substituted or unsubstituted heteroalkyl, a hydroxyl-

protecting group, a phosphate moiety, a phosphodiester moiety, a 7 phosphorus-containing internucleotide bridge of a nucleic acid, a solid 8 support, a carrier molecule and $-OP(O)(OR^{o})(N(R^{p}R^{q}))_{2}$ 9 wherein 10 R^o, R^p and R^q are members independently selected from H, substituted or 11 unsubstituted C₁-C₆ alkyl and substituted or unsubstituted C₁-C₆ 12 13 heteroalkyl; and s is an integer from 1 to 20. 14 (Original) The xanthene dye according to claim 14, wherein R^o is CH₂CH₂CN. 1 15. (Original) The xanthene dye according to claim 14, wherein at least one of R^x and 1 16. 2 R^y comprises a moiety having the formula: ξ —L⁴—R^z 3 wherein 4 L⁴ is a member selected from a bond, substituted or unsubstituted alkyl and 5 substituted or unsubstituted heteroalkyl; and 6 7 R^z is a member selected from a reactive functional group, solid support, a nucleic acid, a saccharide and a peptide. 8 (Original) The xanthene dye according to claim 16, wherein L⁴ comprises a 1 17. 2 moiety having the formula: 3

(Original) The xanthene dye according to claim 1, wherein said carrier molecule

wherein Z³ is a member selected from CH₂ and C=O.

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further comprises a quencher moiety.

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- 1 19. (Original) The xanthene dye according to claim 18, wherein said xanthene dye 2 and said quencher comprise a donor-acceptor energy transfer pair.
 - 20. (Original) The xanthene dye according to claim 18, wherein said quencher has substantially no native fluorescence.
- 21. (Original) The xanthene dye according to claim 20, wherein said quencher comprises at least three residues selected from aryl, substituted aryl, heteroaryl, substituted heteroaryl and combinations thereof, wherein at least two of said residues are covalently linked via an exocyclic diazo bond.
- 1 22. (Original) The xanthene dye according to claim 1, wherein said xanthene dye is 2 attached to a nucleic acid at a position which is a member selected from the 3'-terminus, the 5'-3 terminus, a nucleobase, and a phosphorus-containing internucleotide bridge of said nucleic acid.
- 1 23. (Original) The xanthene dye according to claim 18, wherein said nucleic acid is a 2 probe which is a member selected from molecular beacons, scorpion probes, sunrise probes, 3 conformationally assisted probes and TaqMan™ probes.
- 1 24. (Original) The xanthene dye according to claim 1, wherein said carrier molecule 2 is a peptide comprising a cleavage recognition site for an enzyme.
 - 25. (Original) The xanthene dye according to claim 24, wherein said peptide comprises a cleavage recognition site for a protease.
- 26. (Original) The xanthene dye according to claim 24, wherein said cleavage
 recognition site is for an enzyme selected from trypsin, enterokinase, HIV-1 protease,
 prohormone convertase, interleukin-1b-converting enzyme, adenovirus endopeptidase,
 cytomegalovirus assemblin, leishmanolysin, β-secretase for amyloid precursor protein, thrombin,
 renin, angiotensin-converting enzyme, cathepsin-D and a kininogenase.
 - 27. (Original) The xanthene dye according to claim 1, in which R^{12} has the formula:

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3 wherein

- L^{1a} is a member selected from substituted or unsubstituted alkyl, and substituted or unsubstituted heteroalkyl groups; and
 - R^{2a} and R^{3a} are members independently selected from H, substituted or unsubstituted alkyl, and substituted or unsubstituted heteroalkyl, and R² and R³, together with the nitrogen to which they are attached, are optionally joined to form a ring which is a member selected from substituted or unsubstituted C₅-C₇ cycloalkyl and substituted or unsubstituted 5-7-membered heterocycloalkyl.
 - 28. (Original) The xanthene dye according to claim 27, in which L^{1a} does not comprise a member selected from a carboxylic acid and a carboxylic acid ester.
- 1 **29.** (Original) A method for determining whether a sample contains an enzyme, said 2 method comprising:
 - (a) contacting said sample with a peptide construct comprising:
 - i) a xanthene dye according to claim 1;
 - ii) a quencher; and
- 6 iii) a cleavage recognition site for said enzyme,
- wherein said peptide is in a conformation allowing donor-acceptor energy transfer
 between said fluorophore and said quencher when said fluorophore is
 excited;
 - (b) exciting said xanthene dye; and
 - (c) determining a fluorescence property of said sample, wherein the presence of said enzyme in said sample results in a change in said fluorescence property.

1	30. (Original) A method for determining whether a compound alters an activity of an
2	enzyme, said method comprising:
3	(a) contacting a sample comprising said enzyme and said compound with a peptide
4	construct comprising:
5	i) a xanthene dye according to claim 1;
6	ii) a quencher; and
7	iii) a cleavage recognition site for said enzyme,
8	wherein said peptide is in a conformation allowing donor-acceptor energy transfer
9	between said xanthene dye and said quencher when said xanthene dye is
10	excited;
11	(b) exciting said xanthene dye; and
12	(c) determining a fluorescence property of said sample, wherein said activity of said
13	enzyme in said sample results in a change in said fluorescence property.
1	31. (Original) A method for detecting a nucleic acid target sequence, said method
2	comprising:
3	(a) contacting said target sequence with a detector oligonucleotide comprising a target
4	binding sequence, said detector oligonucleotide having linked thereto,
5	i) a xanthene dye according to claim 1; and
6	ii) a quencher,
7	wherein said detector nucleic acid is in a conformation allowing donor-acceptor
8	energy transfer between said xanthene dye and said quencher when said
9	xanthene dye is excited;
10	(b) hybridizing said target binding sequence to said single-stranded target sequence,
11	thereby altering said conformation of said detector oligonucleotide, causing a change
12	in a fluorescence parameter; and

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- (c) detecting said change in said fluorescence parameter, thereby detecting said nucleic
 acid target sequence.
 - 32. (Original) The method according to claim 31, wherein said complementary strand is synthesized in a target amplification reaction.
 - 33. (Original) The method according to claim 31, wherein said complementary strand is synthesized by extension of the target sequence using said detector oligonucleotide as a template.
- 1 34. (Currently Amended) The method according to claim 31, wherein said 2 fluorescence parameter is detected in [[-]]real-time.
 - **35.** (Original) A method for detecting amplification of a target sequence comprising, in an amplification reaction:
 - (a) hybridizing to said target sequence a detector oligonucleotide comprising a single-stranded target binding sequence and an intramolecularly associated secondary structure 5' to said target binding sequence, wherein at least a portion of said detector sequence is a single stranded tail which is available for hybridization to said target sequence, said detector oligonucleotide having linked thereto,
 - i) a xanthene dye according to claim 1; and
 - ii) a quencher,
 - wherein said detector nucleic acid is in a conformation allowing donor-acceptor energy transfer between said xanthene dye and said quencher when said xanthene dye is excited;
 - (b) extending said hybridized detector oligonucleotide on said target sequence with a polymerase to produce a detector oligonucleotide extension product and separating said detector oligonucleotide extension product from said target sequence;

- (c) hybridizing a primer to said detector oligonucleotide extension product and extending the primer with said polymerase, thereby linearizing said intramolecularly associated secondary structure and producing a change in a fluorescence parameter; and
- (d) detecting said change in said fluorescence parameter, thereby detecting said target sequence.
- **36.** (Original) The method according to claim **35**, wherein said target sequence is amplified by a method selected from Strand Displacement Amplification, Polymerase Chain reaction, Self Sustained Sequence Replication, Transcription Mediated Amplification, and Nucleic Acid Sequence Based Amplification.
- 1 37. (Original) The method according to claim 35, wherein said secondary structure 2 further comprises a partially or entirely single-stranded restriction endonuclease site.
 - 38. (Original) The method according to claim 35, wherein a change in fluorescence intensity is detected.
- 1 **39.** (Original) The method according to claim **38**, wherein said change in fluorescence intensity is detected in real-time.
 - 40. (Original) The method according to claim 35, wherein said intramolecularly associated secondary structure comprises a portion of said target binding sequence.
- 1 41. (Original) A method of preparing a conjugate between a nucleic acid and a 2 xanthene dye having the formula:

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4	in which
5	R ¹ , R ² , R ⁴ , R ⁵ , R ⁶ , R ⁷ , R ⁸ , R ⁹ , R ¹⁰ and R ¹¹ are independently selected from substituted or
6	unsubstituted alkyl, substituted or unsubstituted heteroalkyl, substituted or
7	unsubstituted aryl, substituted or unsubstituted heteroaryl, substituted or
8	unsubstituted heterocycloalkyl, halogen, H, NO ₂ , CN and C(Z ¹)R ¹⁴ , NR ¹⁵ R ¹⁶ and
9	Z^2R^{16} ;
10	R^3 is selected from Z^2R^{16} and $NR^{15}R^{16}$
11	wherein
12	Z ¹ is a member selected from O, S and NH;
13	Z ² is a member selected from O and S;
14	R ¹⁵ is a member selected from H, substituted or unsubstituted alkyl, and
15	substituted or unsubstituted heteroalkyl;
16	R ¹⁶ is selected from H, substituted or unsubstituted alkyl, substituted or
17	unsubstituted heteroalkyl, C(Z ³)R ¹⁷ , and a nitrogen-containing reactive
18	group comprising R ¹⁵ and R ¹⁶ , together with the nitrogen to which they are
19	attached, wherein said reactive group is a member selected from -NHNH ₂ ,
20	-N=C=S and $-N=C=O$
21	wherein
22	Z ³ is a member selected from O, S and NH;
23	R ¹⁷ is a member selected from substituted or unsubstituted alkyl,
24	substituted or unsubstituted heteroalkyl, OR ¹⁸ , and NR ¹⁹ R ²⁰
25	wherein
26	R ¹⁸ is a member selected from H, substituted or unsubstituted
27	alkyl, substituted or unsubstituted heteroalkyl, substituted
28	or unsubstituted aryl, substituted or unsubstituted
29	heteroaryl and C(O)R ²¹
30	wherein

31	R ²¹ is substituted or unsubstituted alkyl or substituted or
32	unsubstituted heteroalkyl;
33	R ¹⁹ and R ²⁰ are members independently selected from H,
34	substituted or unsubstituted alkyl and substituted or
35	unsubstituted heteroalkyl
36	Y is a member selected from $C(O)$ and $S(O)_2$;
37	X is a member selected from (NR ²² R ²³) and (O)
88	wherein
39	R ²² and R ²³ are members independently selected from H, substituted or
10	unsubstituted alkyl and substituted or unsubstituted heteroalkyl; and
11	R ¹² and R ¹³ are members independently selected from substituted or unsubstituted alkyl,
12	substituted or unsubstituted heteroalkyl, substituted or unsubstituted heterocycloalkyl, substituted
13	or unsubstituted aryl and substituted or unsubstituted heteroaryl, with the proviso that at least one
14	of R ¹² or R ¹³ comprises said nucleic acid,
15	said method comprising:
16	(a) contacting a precursor of said conjugate comprising nucleic acid protecting groups
17	with a mixture of amine and alcohol, thereby removing said protecting groups.